## IN THE CLAIMS

- 1-24. (canceled)
- 25. (currently amended) A method of treating a Chlamydia infection in a patient, the method comprises the step of administering to the patient a therapeutically effective amount of an immunogenic protein, wherein the immunogenic protein is <a href="Chlamydia"><u>Chlamydia</u></a> ribosomal protein L7/L12, a homologue of <a href="Chlamydia"><u>Chlamydia</u></a> ribosomal protein L7/L12.
- 26. (currently amended) The method of claim 25 wherein the immunogenic protein is <u>Chlamydia</u> ribosomal protein L7/L12.
- 27. (currently amended) The method of claim 26 wherein the protein has a MW of about 15.8 kDa and a pI of about 4.8 the MW and pI characteristics of protein 12 (as set out in Table II on page 15).
- 28. (currently amended) The method of claim 26 wherein the protein has an N-terminal amino acid sequence of TTESLETLVE (SEQ ID NO:2) disclosed in Table III on page 16.
- 29. (previously presented) The method of claim 25 wherein the protein is a fragment of ribosomal protein L7/L12.
- 30. (canceled)
- 31. (currently amended) The method of claim 29 wherein the fragment comprises at least 7 consecutive amino acids of ribosomal protein L7/L12 the protein.
- 32. (currently amended) The method of claim 25 wherein the immunogenic protein is a homologue of <u>Chlamydia</u> ribosomal protein L7/L12.

- 33. (currently amended) The method of claim 32 wherein the homologue has greater than 50% identity to <u>Chlamydia</u> ribosomal protein L7/L12.
- 34. (currently amended) The method of claim 33 wherein the homologue has greater than 90% identity to <u>Chlamydia</u> ribosomal protein L7/L12.
- 35. (currently amended) The method of claim 32 wherein the homologue has a MW of about 15.8 kDa and a pI of about 4.8 the MW and pI characteristics of protein 12 (as set out in Table II on page 15).
- 36. (canceled)
- 37. (currently amended) A method of preventing a Chlamydia infection in a patient, the method comprises the step of administering to the patient a prophylatically effective amount of an immunogenic protein, wherein the immunogenic protein is <a href="Chlamydia">Chlamydia</a> ribosomal protein L7/L12, a homologue of <a href="Chlamydia">Chlamydia</a> ribosomal protein L7/L12, or a fragment of ribosomal protein L7/L12.
- 38. (currently amended) The method of claim 37 wherein the protein is <u>the Chlamydia</u> ribosomal protein L7/L12.
- 39. (currently amended) The method of claim 38 wherein the protein has a MW of about 15.8 kDa and a pI of about 4.8 the MW and pI characteristics of protein 12 (as set out in Table II on page 15).
- 40. (currently amended) The method of claim 38 wherein the protein has an N-terminal amino acid sequence of TTESLETLVE (SEQ ID NO:2) disclosed in Table III on page 16.
- 41. (previously presented) The method of claim 37 wherein the immunogenic protein is a fragment of ribosomal protein L7/L12.

- 42. (canceled)
- 43. (currently amended) The method of claim 41 wherein the fragment comprises at least 7 consecutive amino acids of <u>ribosomal protein L7/L12</u> the protein.
- 44. (currently amended) The method of claim 37 wherein the immunogenic protein is a homologue of <u>Chlamydia</u> ribosomal protein L7/L12.
- 45. (currently amended) The method of claim 44 wherein the homologue has greater than 50% identity to Chlamydia ribosomal protein L7/L12.
- 46. (currently amended) The method of claim 45 wherein the homologue has greater than 90% identity to <u>Chlamydia</u> ribosomal protein L7/L12.
- 47. (new) The method of claim 44 wherein the homologue has a MW of about 15.8 kDa and a pI of about 4.8 the MW and pI characteristics of protein 12 (as set out in Table II on page 15).
- 48. (canceled)